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APPLICATION NO.	FI	LING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/774,888	02/09/2004		Brett P. Monia	ISIS0074-101 (PTS0009US.C	8872	
27180	7590	09/26/2006		EXAMINER		
ISIS PHARMACEUTICALS INC ZARA, JANE J					JANE J	
1896 RUTH	ERFORD	RD.				
CARLSBAD, CA 92008				ART UNIT	PAPER NUMBER	
				1635		

DATE MAILED: 09/26/2006

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No.	Applicant(s)	
		10/774,888	MONIA ET AL.	
Office Action Summary		Examiner	Art Unit	
		Jane Zara	1635	
Period fo	The MAILING DATE of this communica or Reply	tion appears on the cover sheet	vith the correspondence address	
A SHOWHIC - External after - If NO - Failu Any (	ORTENED STATUTORY PERIOD FOR CHEVER IS LONGER, FROM THE MAINS IN THE M	LING DATE OF THIS COMMUN 37 CFR 1.136(a). In no event, however, may a cation. ory period will apply and will expire SIX (6) MO , by statute, cause the application to become	ICATION.  The reply be timely filed  ONTHS from the mailing date of this communication  ABANDONED (35 U.S.C. § 133).	
Status				
2a)□	Responsive to communication(s) filed of This action is <b>FINAL</b> . 2b) Since this application is in condition for closed in accordance with the practice	☐ This action is non-final.  allowance except for formal ma	·	s
Dispositi	on of Claims			
5)□ 6)□ 7)□ 8)⊠	Claim(s) 1-20 is/are pending in the app 4a) Of the above claim(s) is/are Claim(s) is/are allowed. Claim(s) is/are rejected. Claim(s) is/are objected to. Claim(s) 1-20 are subject to restriction on Papers	withdrawn from consideration.		
10)	The specification is objected to by the E The drawing(s) filed on is/are: a Applicant may not request that any objection Replacement drawing sheet(s) including the The oath or declaration is objected to b	) accepted or b) objected to on to the drawing(s) be held in abeyone correction is required if the drawing	ance. See 37 CFR 1.85(a). g(s) is objected to. See 37 CFR 1.121(	(d).
Priority u	ınder 35 U.S.C. § 119			
a)[	Acknowledgment is made of a claim for All b) Some * c) None of:  1. Certified copies of the priority do  2. Certified copies of the priority do  3. Copies of the certified copies of application from the Internationalise the attached detailed Office action for the certified copies of application from the Internationalise the attached detailed Office action for the certified copies of application from the Internationalise the attached detailed Office action for the certified copies of the certified copies of application from the Internationalise the attached detailed Office action for the certified copies of the priority do  3. Copies of the certified copies of the priority do  3. Copies of the certified copies of the priority do  4. Copies of the certified copies of the priority do  5. Copies of the certified copies of the priority do  6. Copies of the certified copies of the priority do  6. Copies of the certified copies of the priority do  8. Copies of the certified copies of the priority do  9. Copies o	cuments have been received. cuments have been received in the priority documents have bee I Bureau (PCT Rule 17.2(a)).	Application No n received in this National Stage	
2)  Notic 3)  Infor	t(s) e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO nation Disclosure Statement(s) (PTO/SB/08) r No(s)/Mail Date	948) Paper No	Summary (PTO-413) o(s)/Mail Date Informal Patent Application 	

## **DETAILED ACTION**

Claims 1-20 are pending in the instant application.

## Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121: Please elect a single target region from claims 1, 14, 16, 17, 19.

The inventions are distinct, each from the other because of the following reasons:

The different inventions drawn to antisense oligonucleotides that are directed to each target region of SEQ ID NO: 4 are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case, the oligonucleotides and methods comprising them are biologically, structurally and functionally different and distinct from each other. The methods involving the use of a distinct oligonucleotide utilize a different and distinct composition, and so utilize distinct methods steps from each other. For these reasons, the inventions of these different Groups are patentably distinct.

Furthermore, searching the inventions of Groups comprising all of these different oligonucleotide molecules and target regions, and the methods comprising them together would impose a serious search burden. In the instant case, the search of the distinct methods and compositions are not coextensive. There is a search burden also in the non-patent literature. Prior to the concomitant construction and utilization of the

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different nucleic acid constructs of interest there may be journal articles devoted solely to one Group that would not have described the compositions and methods of the other Group. Searching, therefore is not coextensive. As such, it would be burdensome to search the inventions of the different Groups together.

Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the methods comprising administration of different nucleic acid oligonucleotides are unrelated as they comprise distinct steps and utilize different nucleic acid constructs which demonstrates that each method has a different mode of operation. The methodology and materials necessary for each of these distinct methods differ significantly, and each Group constitutes a biologically, chemically and functionally distinct and different composition and method and therefore each involves a patentably distinct invention. Therefore, each method is divergent in materials and steps. For these reasons the inventions of these different Groups are patentably distinct.

Pursuant to 35 U.S.C. 121 and 37 C.F.R. 1.141, the different oligonucleotides and target regions listed in claims 1, 14, 16, 17, 19, and encompassed by claims 1-20 are subject to restriction. In the instant case, one independent and distinct antisense oligonucleotide sequence will be examined in a single application without restriction. Those sequences which are patentably indistinct from the sequence or region selected by the applicant will also be examined.

Claims 1-20 specifically embrace different oligonucleotides with different target regions. Each of these antisense oligonucleotides and corresponding target region is considered to be structurally independent, because each is represented by a unique nucleotide sequence. Furthermore, a search of all the sequences claimed presents an undue burden on the Patent and Trademark Office to search and examine. In view of the foregoing, applicants are required to elect up to 1 (one) antisense oligonucleotide and corresponding target region sequence.

Claims 1 and 14 specifically embrace different oligonucleotides and target regions. Each of these oligonucleotides and corresponding target regions is considered to be structurally independent, because each is represented by a unique nucleotide sequence. Furthermore, a search of all the sequences claimed presents an undue burden on the Patent and Trademark Office to search and examine. In view of the foregoing, applicants are required to elect up to 1 oligonucleotide (SEQ ID No.) and corresponding target sequence.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, restriction for examination purposes as indicated is proper.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one

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or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

## Conclusion

Certain papers related to this application may be submitted to Art Unit 1635 by facsimile transmission. The faxing of such papers must conform with the notices published in the Official Gazette, 1156 OG 61 (November 16, 1993) and 1157 OG 94 (December 28, 1993) (see 37 C.F.R. ' 1.6(d)). The official fax telephone number for the Group is 571-273-8300. NOTE: If Applicant does submit a paper by fax, the original signed copy should be retained by applicant or applicant's representative. NO DUPLICATE COPIES SHOULD BE SUBMITTED so as to avoid the processing of duplicate papers in the Office.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to **Jane Zara** whose telephone number is **(571) 272-0765**. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Peter Paras, can be reached on (571) 272-4517. Any inquiry regarding this application should be directed to the patent analyst, Katrina Turner, whose telephone number is (571) 272-0564. Any inquiry of a general nature or relating to the status of this application should be directed to the Group receptionist whose telephone number is (703) 308-0196.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for Application/Control Number: 10/774,888 Page 6

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published applications may be obtained from either Private PAIR or Public PAIR.

Status information for unpublished applications is available through Private PAIR only.

For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Jane Zara 9-19-06

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